Massachusetts Department of Public Health Massachusetts Immunization Program (MIP)

#### **MODEL STANDING ORDERS**

## **Hepatitis A Vaccine, Inactivated**

These model standing orders are current as of January 2004. They should be reviewed carefully against the most current recommendations and may be revised by the clinician signing them.

# Hepatitis A vaccination is <u>indicated</u> for preexposure protection from hepatitis A virus (HAV) infection for susceptible persons $\geq 2$ years of age in the following groups:

- Persons traveling to, or working in, countries that have high or intermediate endemicity (Asia [excluding Japan], Africa, Central and South America, the Caribbean, Greenland, and Eastern Europe). The vaccine should be administered > 1 month prior to travel.
- Men who have sex with men
- Users of injecting and noninjecting illegal drugs
- Persons who have clotting factor disorders
- Persons who have chronic liver disease
- Persons who have occupational risk for infection (work with HAV-infected primates or with HAV in a research laboratory setting)
- Persons in communities where HAV outbreaks occur (if local epidemiological data indicate it is feasible)
- Children who live in states, counties, or communities with high rates of HAV infection (e.g., the average annual HAV infection rate during 1987-1997 was ≥ 20 cases per 100,000 population, or approximately twice the national average)

# Hepatitis A vaccination may be considered for preexposure protection from HAV infection in susceptible persons $\geq 2$ years of age in the following groups:

- Persons who work in food-service establishments or who are food handlers (if local epidemiological data determine it is cost effective)
- Child care center staff and attendees with ongoing or recurrent outbreaks (if indicated by local epidemiological data)
- Children who live in states, counties or communities with intermediate rates of HAV infection (e.g., the average annual rate of HAV infection during 1987-1997 was 10 – 19 cases per 100,000 population, or approximately the national average
- Any person wishing to obtain immunity

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- 1. Provide patient, parent or legal representative with a copy of the Vaccine Information Statement (VIS) and answer any questions.
- 2. Screen for contraindications according to Table 1.
- 3. Give the correct dose of hepatitis A vaccine intramuscularly (IM) according to the recommended schedule (see Table 2) **Always check the package insert prior to administration of any vaccine**. Administer IM vaccines at a 90° angle with a 22- to 25-gauge needle.
  - a. Toddlers ( $\geq$  2 years of age) and older children: Administer in to the anerolateral aspect of the thigh or deltoid, using a 7/8- to 1½-inch needle, depending on the size of the muscle. For toddlers, you can use the anterolateral thigh, but the needles should be longer, usually 1 inch. The deltoid is preferred for immunization of adolescents.
  - b. Adolescents and young adults (≤ 18 years of age): Administer in the deltoid using a 1- to 2-inch needle, depending on the vaccine recipient's weight (1 inch for females < 70 kg; 1.5 inches for females 70-100 kg; 1 to 1.5 inches for males ≤ 120 kg; and 2 inches for males > 120 kg and females > 100 kg).
  - c. Adults > 18 years of age: Administer in the deltoid using a 1- to 2-inch needle, depending on the vaccine recipient's weight (1 inch for females < 70 kg; 1.5 inches for females 70-100 kg; 1 to 1.5 inches for males ≤ 120 kg; and 2 inches for males > 120 kg and females > 100 kg).
- 4. Administer hepatitis A vaccine simultaneously with all other vaccines indicated according to the recommended schedule and the patient's current vaccine status.
- 5. If possible, observe patient for an allergic reaction for 15 20 minutes after administering vaccine.
- 6. Facilities and personnel should be available for treating immediate hypersensitivity reactions.
- 7. Report clinically significant adverse events to the Vaccine Adverse Event Reporting System (VAERS) at 1-800-822-7967, or via the VAERS website: www.vaers.org.
- 8. Please see the MIP document, *General Protocols for Standing Orders*, for further recommendations and requirements regarding vaccine administration, documentation, and consent.

 Table 1.
 Contraindications and Precautions to Hepatitis A Vaccine

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Clinician's Signature	Date

Valid Contraindications to Hepatitis A Vaccine	Invalid Contraindications (hepatitis A vaccine should be administered)
Anaphylactic reaction to previous dose of	Mild illness with or without low-grade fever
hepatitis A vaccine, alum, 2-phenoxyethanol (Havrix® only)¹, neomycin (Havrix® only), latex (VAQTA® only)², or to any other component of the vaccine (see package insert for specific components)³	Non-anaphylactic allergy to any component for the vaccine
	Local reaction to a previous dose of hepatitis A vaccine
	Immuosuppression
Precautions to Hepatitis A Vaccine:  • Moderate-to-severe acute illness, with or	Personal or family history of non-specific allergies
without fever (temporary precaution) • Pregnancy <sup>4</sup>	Current antimicrobial therapy

<sup>&</sup>lt;sup>1</sup> Persons with a hypersensitivity to 2-phenoxyethanol should receive VAQTA<sup>®</sup>, the Merck preparation, which does **not** contain 2-phenoxyethanol.

 Table 2.
 Recommended Dosages and Schedule for Hepatitis A Vaccine

Age	Dose (Units)		Volume	No.of	Schedule
(Years)	Havrix <sup>®1</sup>	VAQTA <sup>®2</sup>	(mL)	Doses <sup>3,4</sup>	(Months) <sup>4</sup>
2 - 18	720	25	0.5	2	0, 6-12
> 18	1,440	50	1.0	2	0, 6-12

<sup>&</sup>lt;sup>1</sup> Hepatitis A vaccine, inactivated, SmithKline Beecham

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**Travelers** 

<sup>&</sup>lt;sup>2</sup> Persons with hypersensitivity to latex should receive Havrix<sup>®</sup>, the SmithKline Beecham

preparation, which does **not** contain latex.

<sup>3</sup> Persons with a history of anaphylaxis to a vaccine component, but who are at high risk for hepatitis A disease, should be referred to a health care provider for evaluation and possible administration of hepatitis A vaccine.

<sup>&</sup>lt;sup>4</sup> Hepatitis A vaccine should be considered for pregnant women at increased risk for hepatitis A infection.

<sup>&</sup>lt;sup>2</sup> Hepatitis A vaccine, inactivated, Merck & Co., Inc.

<sup>&</sup>lt;sup>3</sup> Travelers should receive the  $1^{st}$  dose  $\geq 4$  weeks prior to travel.

<sup>&</sup>lt;sup>4</sup> Two doses are needed for lasting protection.

### Vaccine

Travelers who are administered the vaccine should receive the first dose  $\geq 4$  weeks prior to travel. Studies show that only 55-60% of vaccine recipients have protective antibodies at 2 weeks post vaccine. Persons traveling to high-risk areas < 4 weeks after the initial dose should be administered IG (see paragraph below), but at a different anatomic injection site. A second dose of vaccine 6-12 months later is necessary for long-term protection.

#### IG

Travelers who are allergic to a vaccine component, whose time prior to departure is < 4 weeks, or who elect not to receive the vaccine, should receive IG.

- If the travel period is < 3 months in duration, travelers should receive a single dose of IG (0.02 mL/kg), which provides protection against hepatitis A for up to 3 months.
- If the travel period is  $\geq$  3 months in duration, travelers should receive IG at 0.06 mL/kg, which provides protection for up to 5 months. At this time, administration of IG must be repeated.

## Pre- and Post-Vaccination Serologic Testing for Susceptibility

- 1. Prevaccination testing of children is not indicated because of their expected low prevalence of infection. Prevaccination may be cost effective when the prevalence of hepatitis A virus infection is > 33%. Persons for whom testing might be indicated include:
- adults who are either born in, or lived for extensive periods in, geographic areas with high endemicity of HAV infection;
- older adolescents and adults in certain population groups (i.e., Native Americans, Alaskan Natives, and Hispanics);
- adults in certain groups that have a high prevalence of infection (see above); and
- adults > 40 years of age.
- 2. Postvaccination testing is *not* indicated because of the high rate of vaccine response among adults and children.

References:		/ /	
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